



Billing Code 4410-09-P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration
Manufacturer of Controlled Substances
Notice of Application
Johnson Matthey Pharmaceutical Materials, Inc.

Pursuant to 21 CFR 1301.33(a), this is notice that on
December 23, 2013, Johnson Matthey Pharmaceutical
Materials, Inc., Pharmaceutical Service, 25 Patton Road,
Devens, Massachusetts 01434, made application by renewal to
the Drug Enforcement Administration (DEA) to be registered
as a bulk manufacturer of the following basic classes of
controlled substances:

Drug	Schedule
Amphetamine (1100)	II
Methylphenidate (1724)	II
Nabilone (7379)	II
Hydrocodone (9193)	II
Alfentanil (9737)	II
Remifentanil (9739)	II
Sufentanil (9740)	II

The company plans to utilize this facility to manufacture small quantities of the listed controlled substances in bulk and to conduct analytical testing in support of the company's primary manufacturing facility in West Deptford, New Jersey. The controlled substances manufactured in bulk at this facility will be distributed to the company's customers.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substances, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODW), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than [INSERT DATE 60 DAYS FROM DATE OF PUBLICATION IN THE FEDERAL REGISTER].

Dated: January 15, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator,
Office of Diversion Control,
Drug Enforcement Administration.

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